US ERA ARCHIVE DOCUMENT

# Product Performance Review By Kevin J. Sweeney, Senior Entomologist

Date: Augus 15, 2006

Dec: 36440<sup>2</sup> )P: 326996

PM: Richard Gebken, PM 10

EPA File Sy bol: 806-GR

Product Nar :

Avon Skin 5 Soft SSS Bug Guard Plus Picaridin Insect Repellent Spray

Active ingre ents: 10% Picaridin

Formulation RTU Pump Spray

Use pattern/ es: Human skin

Request: Ne product that repels mosquitoes, biting midges (no-seeums), sand flies, and

gnats.

OPPTS Guid line: 810.3300 (Studies incorrectly refer to 810.3400)

The followin GLP field studies were submitted to support the subject product

registration:

MRID 4675 005 Evaluation of the Efficacy of a Personal Repellent Against

Mosquitoes

MRID 4675 106 Evaluation of the Efficacy of a Personal Repellent Against

Mosquitoes

MRID 4675 007 Evaluation of the Efficacy of a Personal Repellent Against Biting

Midges

#### Science Reviews of the Submitted Studies

These studies rere also the subject to an ethics review by John Carley.

Note: These soldies are the same as previously reviewed MRIDs for 806-GN

## MRID 46751 05 Evaluation of the Efficacy of a Personal Repellent Against Mosquitoes

The subject p duct formulation was identified as 1004024-010(B) in this study.

Location: The study was conducted in the coastal plain of Georgia, USA. The site was located at the selection was avanuah Canal Museum and Nature Center Site in Savannah, GA. Site ased upon prevailing populations of mosquitoes landing at the rate of 1 to in 250 square cm of exposed skin. Recording sites were rotated during the essure changed. The mosquito species prevalent across these sites was ox. Other species includes Aedes and Ochlerotatus spp.

### Study design

Fifteen subjects served as test subjects and two subjects served as negative control subjects. Two subjects were alternates in the event subjects leave the test. A positive control was not tested.

The repellent ormulation application was made by a syringe to an exposed forearm and lower leg at a olume of 0.47 ml/250 sq. cm. (treatment rate of 1.67 mg/cm<sup>2</sup>). The rest of covered with clothing. Shoes were treated with a permethrin-based vent tick bites. Treatment was made early enough to allow peak mosquito o coincide with the eight hour exposure period. The treated subjects were exposed cont exposed an u rate existed a reated leg for five minutes every hour to determine if an adequate biting is site. The test lasted for eight hours.

**Results:** The spellent was effective for 8 hours on all 30 limbs tested. The repellent did not fail on an of the treated subjects during the eight-hour exposure period. Biting pressure was throughout the testing period, averaging 12.6 to 14.7 bites study.

Conclusion: he data supports a claim of "Repels mosquitoes for up to 8 hours".

### MRID 4675. 06 Evaluation of the Efficacy of a Personal Repellent Against Mosquitoes

The subject resolute formulation was identified as 1004024-010(B) in this study.

Location: 7 e study was conducted in the State of Maine, USA. The site was located at sland, Lake Nicatous, Maine. Site selection was based upon prevailing populations of mosquitoes landing at the rate of 1 to 10 per minute on 250 sq cm of Recording sites were rotated during the day as biting pressure changed. Species prevalent across these sites was Ochlerotatus intrudens.

### Study desig :

Ten subject: erved as test subjects and two subjects served as negative control subjects. Two subject were alternates in the event subjects leave the test. A positive control was not tested.

The repeller formulation application was made by a syringe to an exposed forearm and wer leg on each subject at the rate of 0.47 ml/250 sq. cm. (treatment rate of 1.67 mg/cm.

Treatment was made early enough to allow peak mosquito biting activity to coincide to the eight hour exposure period. The rest of the subject was covered with es were treated with a permethrin-based repellent to prevent tick bites. The treated subject to swere exposed continuously until the First Confirmed Bite (FCB).

Negative condeted to an exposed forearm and wer leg on each subject at the rate of 0.47 ml/250 sq. cm. (treatment rate of 0.47 ml/250 sq. cm.)

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**Results:** The average repellency time based upon the FCB bite test was 7 hrs and 54 minutes with a standard deviation of  $\pm$  24 minutes. The repellent was effective for up to eight hours of 19/20 limbs tested. On the forearm of one subject, the repellent failed at 6 ninutes. Biting pressure was adequate throughout the testing period, averaging 3: o 37 bites throughout the study.

Conclusion The results support a label claim of "Repels mosquitoes for up to 8 hours".

### MRID 4675 107 Evaluation of the Efficacy of a Personal Repellent Against Biting Midges

The subject oduct formulation was identified as 1004024-010(B) in this study.

Location: T study was conducted in the coastal plain of Florida, USA. The sites were located at Coastal plain of Florida. Site selection was based upon prevailing biting midges landing at the rate of 1 to 5 per minute. Recording sites uring the day as biting pressure changed. The biting midge species so these sites was Culicodes furens (poey). Some Culicodes barbosi (Wirth vere also collected.

#### Study desig

Ten subjects erved as test subjects and two subjects served as negative control subjects. Two subjects vere alternates in the event subjects leave the test. A positive control was not tested.

Each subject /as treated up to eight hours before exposure to midges in order to test the repellent at t time of peak biting midge activity. The repellent formulation was applied to the exposed forearm at the volume of approximately 0.47 ml/250 sq. cm. to yield 67 mg/cm². The rest of the subject was covered with clothing. Shoes were reated with exposed con exposed and treated arm for five minutes every hour to determine if an adequate biting rate existed: a site. The test lasted for up to eight hours. Testing was conducted on two consecutive

Results: The epellent was effective for up to eight hours (one subject) against the Culicoides b ng midge species. Repellency duration averaged 4 hours and 18 minutes to minutes with the average of both sessions equal to 5 hours and 48 minutes biting pressure was adequate throughout the testing period.

Conclusion: 'he data support a label claim of "Repels biting midges for up to 6 hours".

### Entomologie s Recommendations:

- 1. The bject studies are acceptable and support the following repellency dura on label claims:
  - a "Repels mosquitoes for up to 8 hours." Reapply after 8 hours
  - b "Repels biting midges (no-secums) for up to 6 hours." The registrant requested a reapplication interval of 5 hours and this reapplication interval is acceptable.
- 2. Data nould be submitted or cited to support the claim for sand flies and gnate black flies) or these pests should be removed from the label.
- 3. Pleas explain how the extended duration and time-release technology is supp ted.
- 4. Pleas remove all references to an invisible, hidden, or unseen barriers for protection.
- 5. The dim for repelling WNV vectors is not supported by the submitted data. Pleas cite or submit additional data with WNV vectors or remove this claim. The disquitoes tested in these studies are not WNV vectors. Species from the general Culex should be tested. Testing should also be conducted against Aede. Albopictus.
- 6. Rem e the claim that skin-so-soft is an antidote to mosquitoes. This is an unsu ported medical claim.